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Horizons in Medicine

## **Valproate: life-saving, life-changing**

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## **Abstract**

Anti-epileptic medications, and valproate principally, are commonly prescribed teratogens. There is significant concern that we are not doing enough to educate clinicians and potential parents about the risks of valproate in pregnancy. There is clear advice from the Medicines and Healthcare products Regulatory Agency (MHRA) and the International League Against Epilepsy (ILAE) about the risks of valproate exposure in utero. Reviews and guidelines that are focussed on foetal risk however fall short in being able to fully replicate the complexity of a real clinical decision. Valproate is certainly life-changing if your child is one of the 10% with a major malformation or 30-40% with a neurodevelopmental disorder, but valproate is also potentially life-saving in the context of ensuring the best possible seizure control for some mothers with epilepsy. There are significant knowledge gaps regarding the risks to mothers who elect to take another drug, or to mother and baby if she comes off medication entirely. We also should be doing more to reduce rates of sudden unexpected death in epilepsy (SUDEP) which is recognised as a key target when evaluating all maternal deaths.

## Introduction

There is a common clinical paradox when selecting the correct treatment for a young woman with genetic generalised epilepsy. On one hand, you can reassure her that about 4 in 5 people will become seizure free, but on the other hand you know that the drug most likely to bring this about has significant deleterious effects. Valproate is a drug that can be life transforming if it controls your epilepsy, but it is highly controversial because of the high rates of neurodevelopmental problems associated with children exposed to it in utero. This clinical conundrum is even more acute when all other drugs have failed to yield adequate seizure control and the only efficacious option that remains is valproate.

Thalidomide has become a touchstone for both the scientific and lay community when describing the potential scale of a drug-side effect problem. Just alluding to the drug 'thalidomide' conjures up both the widespread harm caused to an estimated 10,000 children world-wide, but also the battle to recognise this risk and restrict its use. The drug had and has many important medical uses, but when taken in early pregnancy to combat hyperemesis, affected children were born with significant limb deficits. Although withdrawn in the UK in 1961 it took a long campaign and press-support before there was a compensation settlement. Therefore, when *The Daily Telegraph* headline on 30<sup>th</sup> July 2009 stated that valproate was responsible for 'Worst child poisoning case since thalidomide' – the connotation was stark and emotional (figure 1). I will discuss the many reasons why the thalidomide comparison is understandable, but also why the circumstances are significantly different. This will be with the specific aim of promoting the unanimous advice about valproate in pregnancy from many august agencies, while also recognising that there are important evidence gaps.



**Figure 1:** British newspapers respond to parental concern regarding valproate prescription in pregnancy

### Valproate exposure in utero

Valproate is a recognised teratogen and the major cause of foetal anticonvulsant syndrome. An estimated 10.7% of children exposed to valproate in utero are born with a major congenital malformation (that is; a structural abnormality that is recognisable at birth) in comparison to an estimated 2% of un-exposed infants.[1] This produces the first two problems: 1) doctors are often poor at explaining risk; 2) patients are inexperienced in balancing these risks. This is exacerbated when the research evidence produces population risks '*a certain percent of affected children..*' whereas what we need is individualised risk '*your chance of having an affected baby is..*'. Equally, our prior experiences as physician and parent will sway the way we deal with perceived risk. The second issue is this – the fact that the concern regarding major malformations and valproate is so well known it has blinded us to the new scandal, the new relationship. We, as adult physicians have had 'change blindness' – we have failed to see that the threat of valproate is no longer limited to malformations at birth – but more far-reaching and life-changing developmental issues.

The easiest associations to recognise are when infrequent exposures (unusual drug x) causes a dramatic, instant and infrequently occurring outcome. A much more challenging situation is when a drug which has been available since the late 1960s, such as valproate, produces a delayed and variable effect, such as poorer cognition. Recognition is even harder when the effect is in the patients' offspring. This in part explains why the epilepsy community was so slow to recognise the extent of the risks.

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### The evidence

The mainstay of evidence is epidemiological and as a result we can draw correlations but cannot speak directly to causation. These sources are either large retrospective cohort studies or they are smaller prospective studies. The pregnancy registries have been a great asset. Since 1996 The UK and Ireland Epilepsy and Pregnancy Register ([www.epilepsyandpregnancy.co.uk](http://www.epilepsyandpregnancy.co.uk)) has been collecting information on women with epilepsy who became pregnant, with a view to determining the effect of anti-epileptic drug exposure on the developing foetus. They currently have information on over 10,000 pregnancies and continue to collect around 800 new cases each year; about 25% of all eligible pregnancies. Typically, a mother with epilepsy will contact the register and complete a questionnaire about their

**Commented [RT2]:** Does the mother contact the register herself – does she need an invitation?

health, their epilepsy and their medication – before the outcome of the pregnancy is known. Although there are undoubtedly biases about which types of women are motivated to participate and the ability of researchers to fully identify malformations or developmental issues without examining each child, the scale and rigour of the registries mean that they yield comparable results to each other.

The current concern is about adverse neurodevelopmental outcomes; intellectual impairment and autism. Prospective evidence that intellectual impairment could be identified in children as young as three came in 2009 as the **NEAD** study group published their landmark paper.[2] At this stage the threats of anti-epileptic drugs were feared but not known – the relevant consensus guidelines by the American Academy of Neurology,[3] the American College of Obstetricians and Gynecologists,[4] and the International League against Epilepsy[5] did not differentiate between the anti-epileptic drugs. The authors identified 309 children exposed to anti-epileptic drugs in utero, 258 were tested at 2 or 3 years using standardised age-appropriate tests. 53 were exposed to valproate, 73 to carbamazepine, 84 to lamotrigine and 48 to phenytoin. The mean IQ of the children exposed to valproate was 92 (CI 88-97) which is significantly lower than the IQs of the other groups of children (98(CI 95-102), 101(CI 98-104) and 99(CI 94-104) respectively). This relationship was even more stark when the authors looked at smaller groups – high dose v low dose; the 22 children exposed to high dose valproate had a mean IQ of 87. Higher maternal IQs were related to higher child IQs (as expected) – except in the case of valproate. As expected, some children exposed to valproate had congenital malformations, and lower IQ – but these children alone did not explain the IQ differences. As expected this relationship was confirmed when the study group were retested at 6 years.[6] They helpfully demonstrated that breastfeeding did not increase the risks and conferred the expected benefits to infants' IQ.[7]

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A Cochrane review of 22 observational prospective cohort studies and six register studies confirmed the observation of poorer neurodevelopmental outcomes of pregnancies exposed to valproate.[8] The effect is dose related and exacerbated by polytherapy. In 2015 the **EURAP** study group (an International registry of anti-epileptic drugs and pregnancy) published on 1224 pregnancies where valproate was taken alone and a further 364 where it was taken alongside another epilepsy drug. The rate of congenital malformation was 10% in children exposed to valproate (11.3% if exposed to valproate and lamotrigine, 11.7% exposed to valproate and another antiepileptic drug that wasn't lamotrigine). However, the dose really mattered; when more than 1.5g of valproate was used these rates increased to 24% for valproate alone; 31% for valproate and lamotrigine; and 19.2% valproate and another drug.[9]

**Commented [RT4]:** Define acronym

In 2013 a Danish population based study of 655,615 children born 1996-2006 identified that mothers

who took valproate in utero were five times more likely to have a child diagnosed with autism. This equates to a rate of 4.4/100 risk of autism spectrum disorder and a 2.5/100 risk of autism in childhood.[10] Well-designed prospective studies have delineated the autistic traits in children.[11]

### **Advice**

In January 2015 the Medicines and Healthcare Products Regulatory Agency (UK) took the important step of issuing a statement regarding the safety of valproate when taken in pregnancy.[12,13] This advice followed on from the European Medicines Agency in 2014 who recommended strengthening the restrictions on the use of valproate in women and girls. Physicians were quicker to release statements than governmental agencies; in 2009 the American Academy of Neurology recommended that valproate should be avoided in pregnancy whenever possible.[14]

The MHRA stated “No-one should stop taking valproate without discussing it first with their doctor and the benefits of valproate treatment must be carefully balanced against the risks.” This message was echoed by the French Health Minister, Marisol Touraine, earlier this year, who announced practical measures such as adding an illustration to the verbal warning on the medication-packaging, and altering doctors' prescription software to include targeted information. The joint task force of the International League Against Epilepsy commission on European affairs and the European Academy of Neurology responded with a detailed letter.[15] Their advice included that women of child-bearing potential must use effective contraception during treatment.

The apparently contrary advice is that women should remain on medication throughout pregnancy. This oft repeated edict is based, primarily, on the risk of convulsions on the health of the foetus. Although this intuitively feels like the safest advice, it has never been studied with sufficient rigor and our evidence primarily comes from extrapolation from studies of eclampsia (where the pathophysiology of the seizures is very different) and from experiments on rats. Papers attempting to identify the risks of seizures on gestational outcome are all limited by the fact that there is often something inherently different about women with epilepsy who either seize frequently or need multiple drugs to control their epilepsy. A Thai study of 1016 women with epilepsy identified that 49.5% had seizures in pregnancy. Women with epilepsy who had seizures during pregnancy were more likely to have lower educational achievement, to be unmarried, have a lower family income and their male partners were younger.[16] A study of 106 women with epilepsy on medication showed that babies of women who had one or more seizure in pregnancy were more likely to have a shorter gestational age, and a five-times higher risk of being born before term.[17] However the same study demonstrated that there were drug risks too – women taking two or more anti-epilepsy drugs were

more likely to have small for gestational age babies.

There is a compulsory medical Birth registry in Norway and a study of 1900 women with epilepsy (66% on no medication) identified that the majority of the poorer outcomes were seen in women who had epilepsy significant enough to need drug control. Both groups (medication v. no medication) exhibited seizures in the neonatal period. There were perinatal deaths in both groups (0.6% v 0.7%) which was non-significant.[18] The advice that women should remain on their medication therefore does not equate that *all* women with epilepsy should be on medication in pregnancy. A UK study demonstrated that when pregnant, women with epilepsy were **twice as likely** to come off their medication – but should this have happened prior to conception?[19] We are currently unable to tell which women would be safe to trial a withdrawal of medication when offering pre-conception counselling.

**Commented [RT5]:** Twice as likely as who?

### Genetic generalised epilepsy

The majority of mothers with epilepsy who take valproate have GGE (genetic generalised epilepsy); the evidence for the genetic component is compelling and there is an increased risk of epilepsy in first-degree relatives. The genetic architecture is complex and not yet fully elucidated.[20,21] Clearly the maternal 'epilepsy genes' plus any paternal contribution to risk can and will be transmitted to the infant. It is unknown whether parental genomic factors contribute towards valproate associated neuro developmental disorders. Are women with GGE, therefore, at greater risk of neurodevelopmental disorders in their children than women with a focal epilepsy? This is very hard to unpick because current practice is that valproate is the primary drug for GGE. GGE is the only syndrome in which valproate could be justified in pregnancy, due to our understating of the risks. Many pregnancy registers and epidemiological studies have been unable to report maternal epilepsy syndrome accurately. A study of 248 valproate exposed pregnancies in 2008 identified that 126 had idiopathic generalised epilepsy and of these, 15 had a major malformation. They did not report on neurodevelopmental outcomes. Reports from other disorders are scanty – not least because women with migraine and bipolar tend to come off medication before pregnancy.[19]

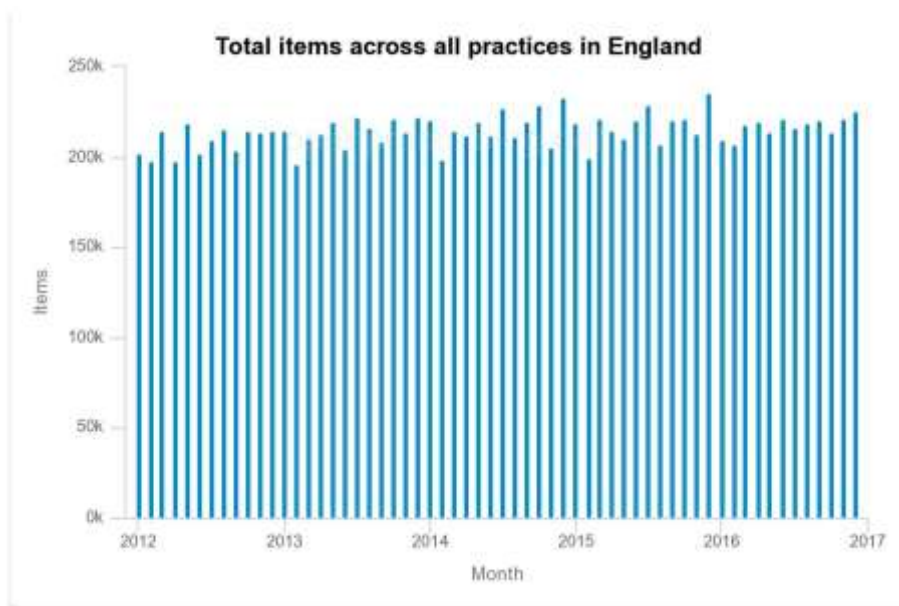
### Scale of the problem

One of the reasons why this is a bigger issue than thalidomide is that valproate in Europe and across



the world is much more frequently prescribed. Women with epilepsy have 1% of all pregnancies and valproate has been widely prescribed since the late 1960s.[22] It is estimated, therefore, that there are thousands of people in the UK living with iatrogenic valproate associated neurodevelopmental disorders. Valproate remains a commonly prescribed drug and in 2002 was the most commonly prescribed antiepileptic drug worldwide.[23] We know that there has been a decade long reduction in valproate prescription in women aged 14-45 years in Wales.[24] Valproate in 2000/2001 was the most prescribed drug to these women (45%) falling to 24% by 2009/10. Younger women (18 years and under) also saw valproate prescriptions fall from 74% to 46% during this period. **Causing more concern** however, current UK data (figure 2) demonstrates that there has been no overall shift in the number of units of valproate that are being prescribed in primary care from 2012 to date. There is a significantly sized persistent group that remain on the drug despite strong evidence and clear advice, why might this be?

**Commented [RT6]:** Meaning of sentence difficult to understand – “more” doesn’t seem to match with falling rates of Valproate use. Also need to draw attention to the fact that the same number of units being sold would seem to contradict that the rates of prescription have fallen.



**Figure 2.** Total Number of items of sodium valproate prescribed by GP practices across England from January 2012 to January 2017. [www.OpenPrescribing.net](http://www.OpenPrescribing.net) EBM DataLab, University of Oxford, 2017[43]

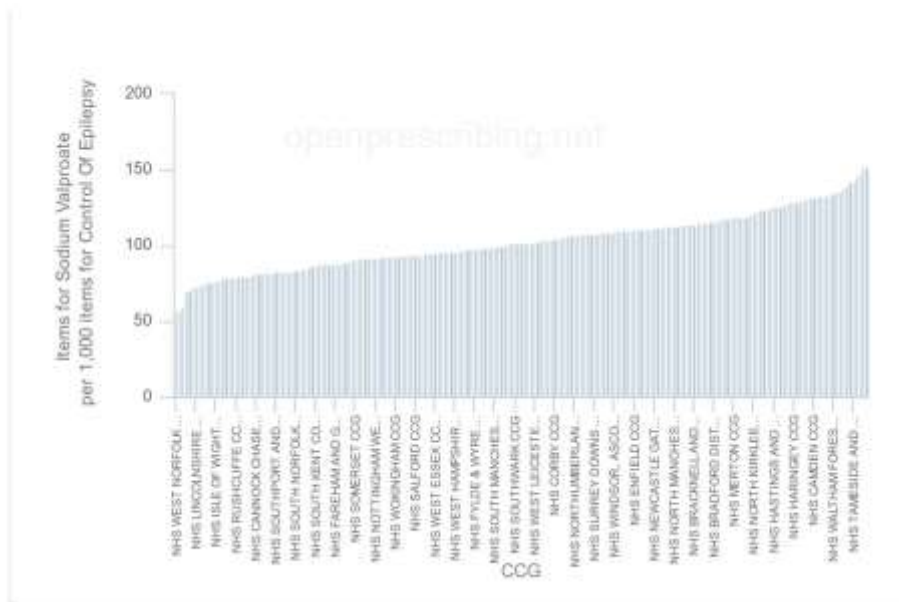
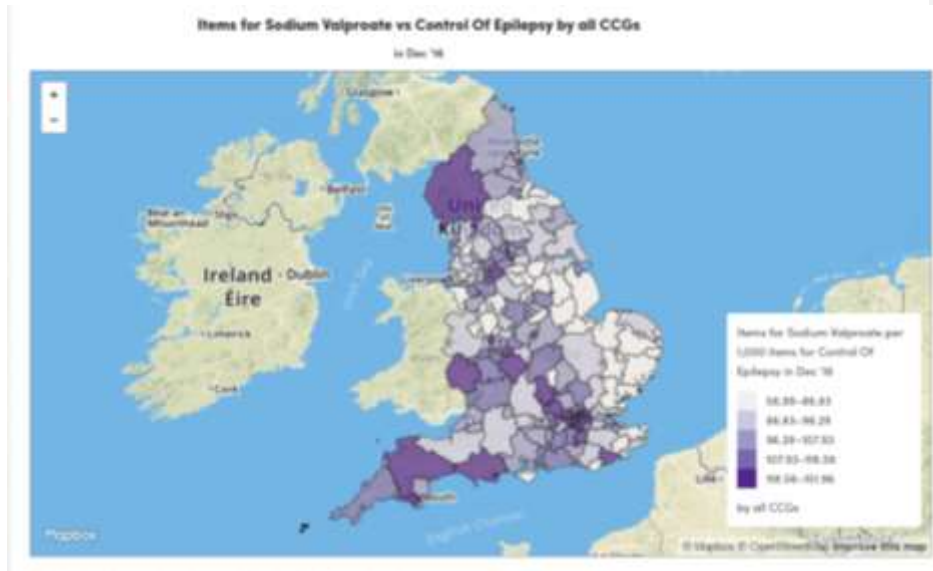


Figure 3. The graph shows the proportion of valproate items prescribed compared to all items prescribed for control of epilepsy. All CCGs are shown on the graph online. (www.OpenPrescribing.net, EBM DataLab, University of Oxford, 2017)[43]

Figure 3 shows the variety in the prescription of valproate across CCGs in England compared to all drugs for the control of epilepsy. CCGs such as West Norfolk (far left) in December 2016 used 560 items of valproate vs 9827 items for epilepsy in comparison to NHS Stockport (far right) which used 2151 items of valproate compared to 14155 items for epilepsy control. This equates to a 2.6 fold difference across the UK (57 items of epilepsy per 1000 items to control epilepsy vs 152). Does the presence of a strong epilepsy community as indicated by a local neurosciences centre influence valproate prescription? (Figure 4). Epilepsy prevalence is highly correlated with levels of deprivation[25] – however the map in figure 4 is not a 'deprivation map' of England. CCGs in the top 20% should embrace the challenge of reducing their valproate prescribing and learn how this is possible from the best performing CCGs.

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**Figure 4.** A map showing the data from figure 3 – the geographical variability of valproate prescribing compared to all items for epilepsy. (www.OpenPrescribing.net, EBM DataLab, University of Oxford, 2017)[43] There appears to be an East v West divide. Some CCG areas of higher valproate prescription – do not have a neurosciences centre: NHS Cumbria, Herefordshire, South Tyneside, North Kirklees. There is a 'M40-M25' corridor of high prescription around Oxford and North and West London.

#### Valproate is an effective drug

One of the reasons the main reason that valproate is not like thalidomide is that we cannot simply stop prescribing it completely. Influential, UK-based, pragmatic clinical trials such as **SANAD** reinforce the clinical view that valproate is superior to many of the subsequent anti-epileptic drugs and is the most efficacious in genetic generalised epilepsy.[26] 716 patients were randomised to open-label lamotrigine, topiramate or valproate. Valproate was superior in terms time to treatment failure (a compound metric of 'do people remain on the drug?' combining both side effects and seizure control). Valproate's strength was primarily in seizure control.

Seizure control is not the only thing important to potential mothers with epilepsy – but it *is* important. Most quality of life outcomes correlate with seizure control; sleep, mood, anxiety, confidence, pain; all are worse when seizures are frequent. UK driving restrictions apply to almost all people with recurrent seizures, meaning that a mother with an absence seizure following a period of control must

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surrender her licence for at least a year before a return to driving. This clearly has a negative consequence on employment and employability despite legislation to protect against overt discrimination.

Furthermore, seizures themselves are not benign. The MBRRACE-UK (Mothers and Babies: Reducing Risk through audits and confidential enquiries across the UK) report in December 2014 recognised that “Epilepsy remains a high risk condition in pregnancy” continuing “multi-agency evidence-based guidelines are urgently required to standardise and improve the care of pregnant and postpartum women with epilepsy.” Epilepsy was a co-morbidity in 17 (7.8%) maternal deaths in pregnancy. Of the women who died in pregnancy not one was reported as taking valproate. Lamotrigine was the most commonly prescribed drug; the pharmacodynamics of lamotrigine mean that some clinicians regularly test levels in pregnancy, whereas many do not.[27] Fourteen deaths are reported on in detail, all are caused by poor seizure control; twelve were SUDEP (sudden unexpected death in epilepsy ) and two were drownings. [28] Only two of these twelve who died from SUDEP had adequate seizure control prior to pregnancy and poor seizure control correlates with an increase in SUDEP risk. Furthermore a review of similar deaths over thirty years concluded that, between 1979-2008, although maternal deaths have decreased, the proportion related to epilepsy have increased.[29] Put bluntly, restricting efficacious drugs from women increases their risk of dying from their epilepsy. Even when a woman is not pregnant, epilepsy confers a twenty-fold increase in death because of SUDEP. SUDEP kills more Britons per year than asthma, or AIDS, or house fires. Hence when the International League Against Epilepsy issued their report in 2015, there were authors rebutting their advice with letters such as “Valproate still has a place in women with epilepsy”.[30]

### **Valproate the drug**

Valproate was not the output of any rational drug discovery programme, rather a serendipitous accident. Pierre Eymard was performing experiments on rat-models of epilepsy in France in the early 1960s. Valproic acid was used as a lipophilic vehicle for compounds that were water-insoluble. Each compound dissolved in the valproic acid was a very acceptable anti-epileptic agent, but so too was the control vehicle with no ‘active’ compound in it. Very quickly they moved to use sodium valproate in rabbits and then other groups performed the first in man studies in 1964. By 1967 it was marketed in France as Depakine – within five years of its discovery.[31]

Valproic acid has several formulations and many indications including migraine prophylaxis, mood-stabilisation and to a lesser degree pain control. This means that advice such as the MHRA advice needs

to be advertised widely beyond the narrow scope of neurology in secondary care. Valproic acid is theorised to have many modes of action: inhibition of GABA transaminase; suppression of repetitive neuronal firing through inhibition of voltage-sensitive sodium channels; it also has a histone deacetylase inhibitor function. The combination of a 'happy accident' discovery and a varied mode of action has stymied researchers as they have sought to identify what is it specifically about the molecule which confers the risk in pregnancy as they aim to design safer valproates.

There are a number of possible and potentially overlapping explanations for the association between exposure to valproate in utero and neurodevelopmental disorders in children. These include: 1) There is a direct effect on brain development; 2) The association is genetic but pleiotropic, driven by genetic overlap between the maternal epilepsy and causes of neurodevelopmental disorder; 3) There is a 'gene x environment' interaction and that a high genetic risk of genetic generalised epilepsy from the mother renders the developing child more susceptible to valproate's effects on brain development; 4) Valproate is potentially mutagenic, and that there is an excess of de novo mutation in the children exposed to valproate.

What is telling is that one of the methods of modelling 'autism' in rats is to expose them to high doses of valproate in utero.[32] Rat models provide a better method of studying the direct brain effects of valproate.[33] Initially exposed rat pups show alteration in medial prefrontal cortex synaptic connectivity. Their principal neurons show decreased excitability but increased local connectivity, due in part to an up-regulation of NMDA receptor expression. The synaptic abnormalities in exposed rats persist into adulthood.

Valproate is primarily used in the epilepsy syndrome genetic generalised epilepsy (synonymous with idiopathic generalised epilepsy).[26] This is a developmental epilepsy syndrome which often starts in childhood or teenage years and can be exacerbated by sleep-deprivation and alcohol. There are also neurocognitive problems associated which include some executive function difficulties and impulsivity problems.[34,35] This is partly why the cognitive outcomes in the offspring in this group were not recognised; in part maternal under-reporting and in part a paternalistic doctorly view that these borderline IQs may be commensurate with maternal abilities. However, in fairness, it is worth adding that experts were asking questions about the safety of epilepsy drugs in pregnancy as early as 1964 – prior to the introduction of valproate.[36] What we have not proven with sufficiently powered studies is the role that maternal epilepsy syndrome plays on the likelihood of neurodevelopmental outcomes in the child.[37] The reasons why this is crucial are laid out below.

### Epilepsy is different

In 2011, Phil Smith and I wrote that 'Epilepsy is different';[38] this Editorial was a call to re-examine the series of contradictions that surround epilepsy – such as: it is potentially invisible, yet stigmatising; controllable, yet a cause of sudden death; common, but so little is known about it by lay audiences. Another contradiction is that family studies and twin registries tell us that it should be highly heritable and yet we know surprisingly little about the genetic architecture of common epilepsies.[20] That transmissible genetic factors are important is incontrovertible and a recent paper identifies for the first time that these mutations are also those that are seen to cause severe childhood epilepsies with autism and intellectual disability.[39]

There are several important research questions that need to be answered

1. Why are some children preferentially affected and can this be determined by studying mothers – ideally prior to conception? Could we identify 'high-risk' and 'low-risk' women to personalise pre-conception counselling?
2. Is valproate the only drug that is working in this way? Are there other drugs working via similar mechanisms that have a similar result but to a lesser degree and so are yet to be discovered?
3. With the understanding now about very rare gene changes in people with common epilepsies; what role does the maternal epilepsy syndrome have on the child? Do women with migraine and bipolar have similar problems?
4. Does valproate cause new mutations in the child? We know that these de novo mutations are an important cause of intellectual disability and autism. Approximately 40% of children with intellectual disability have a causative de novo mutation identifiable with genome sequencing.[39,40] The mechanisms by which any teratogen affects organogenesis are still unclear but there is a large overlap between mutagenicity and teratogenicity. As the evidence that valproate is a teratogen is indisputable, we have to consider whether it is mutagenic.[41] Valproate readily crosses the placenta and circulates in the embryo in higher concentrations than in maternal blood.

### Conclusions

It is clear from consensus statements and an independent review of the evidence that valproate confers a risk of neurodevelopmental disorder to 10% of exposed infants. This risk and this rate is unacceptable to most mothers and when adequate pre-conception counselling is provided they will choose to withdraw from valproate and / or swap valproate for another drug. For women who have

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frequent seizures the smallest dose of valproate that controls their seizures is best and polytherapy should be avoided (particularly combinations of valproate and topiramate).[42] The mother should be central in our decision making process and we should support her if, when appraised of all the evidence, she makes a decision to remain on valproate in pregnancy. She should not be made to feel selfish for considering maternal quality of life factors when making a decision. Tools for decision making such as the [optiongrids.org](http://optiongrids.org) are helpful when weighing up difficult choices such as these.

We need to know how valproate causes neurodevelopmental disorders as the drug remains an important treatment for women with epilepsy. What is the risk to women with epilepsy and their children when they are unmedicated in pregnancy? When does the risk balance tip so that future advice becomes – remain on your drug, or if it is valproate – swap it if you can? In addition to the evidence gaps discussed above, I want to ask: has the time come for it to be compulsory to enrol all women who take valproate in to a national registry? Should we be formally consenting all women aged 14-45 who choose to take valproate, and should there be a national standard **for this?**

**Commented [RT11]:** Fab ☐! However, does it need a punchier sentence to end it – even perhaps: everything points to the answers being “yes” and “yes” “yes”.





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## References

1. Meador K, Reynolds MW, Crean S, Fahrbach K, Probst C Pregnancy outcomes in women with epilepsy: a systematic review and meta-analysis of published pregnancy registries and cohorts. *Epilepsy Res.* 2008; 81(1):1-13.
2. Meador KJ, Baker GA, Browning N, Clayton-Smith J, Combs-Cantrell DT, Cohen M, Kalayjian LA, Kanner A, Liporace JD, Pennell PB, Privitera M, Loring DW, NEAD Study Group. Cognitive function at 3 years of age after fetal exposure to antiepileptic drugs. *N Engl J Med.* 2009; 360(16):1597-605.
3. Harden CL, Hopp J, Ting TY et al. Practice parameter update: management issues for women with epilepsy--focus on pregnancy (an evidence-based review): obstetrical complications and change in seizure frequency: report of the Quality Standards Subcommittee and Therapeutics and Technology Assessment Subcommittee of the American Academy of Neurology and American Epilepsy Society. *Neurology.* 2009;73(2):126-32.
4. <http://www.acog.org/Patients/FAQs/Seizure-Disorders-in-Pregnancy-ACOG-April-24th-2017>
5. Harden, CL, Meador, KJ, Pennell, PB et al. Management issues for women with epilepsy—Focus on pregnancy (an evidence-based review): II. Teratogenesis and perinatal outcomes. *Epilepsia* 2009;50:1237–1246.
6. Meador KJ, Baker GA, Browning N, Cohen MJ, Bromley RL, Clayton-Smith J, Kalayjian LA, Kanner A, Liporace JD, Pennell PB, Privitera M, Loring DW; NEAD Study Group. Fetal antiepileptic drug exposure and cognitive outcomes at age 6 years (NEAD study): a prospective observational study. *Lancet Neurol.* 2013;12(3):244-52.
7. Meador KJ, Baker GA, Browning N et al. Breastfeeding in children of women taking antiepileptic drugs: cognitive outcomes at age 6 years. *JAMA Pediatr.* 2014;168(8):729-36.
8. Adab N, Tudur Smith C, Vinten J, Williamson PR, Winterbottom JB, McKay AJ, Bromley R. Common antiepileptic drugs in pregnancy in women with epilepsy. *Cochrane Database Syst Rev.* 2015;(12):CD004848.
9. Tomson T, Battino D, Bonizzoni E, Craig J, Lindhout D, Perucca E, Sabers , Thomas SV, Vajda F; EURAP Study Group. et al. Dose-dependent teratogenicity of valproate in mono- and polytherapy: an observational study. *Neurology* 2015;85(10):866-72.
10. Christensen J, Grønborg TK, Sørensen MJ, et al. Prenatal valproate exposure and risk of autism spectrum disorders and childhood autism. *JAMA* 2013;309:1696-703.
11. Wood AG, Nadebaum C, Anderson V, Reutens D, Barton S, O'Brien TJ, Vajda F. Prospective assessment of autism traits in children exposed to antiepileptic drugs during pregnancy. *Epilepsia.* 2015;56(7):1047-55.
12. MHRA Drug Safety Update: Medicines related to valproate: risk of abnormal pregnancy outcomes, 22 Jan 2015 [www.gov.uk/drug-safety-update/medicines-related-to-valproate-risk-of-abnormal-pregnancy-outcomes](http://www.gov.uk/drug-safety-update/medicines-related-to-valproate-risk-of-abnormal-pregnancy-outcomes)
13. MHRA Drug Safety Update: Valproate and of risk of abnormal pregnancy outcomes: new communication materials 17 Feb 2016 [www.gov.uk/drug-safety-update/valproate-and-of-risk-of-abnormal-pregnancy-outcomes-new-communication-materials](http://www.gov.uk/drug-safety-update/valproate-and-of-risk-of-abnormal-pregnancy-outcomes-new-communication-materials)
14. Harden CL, Meador KJ, Pennell PB, et al. American Academy of Neurology. American Epilepsy Society Practice parameter update: management issues for women with

- epilepsy—focus on pregnancy (an evidence-based review): teratogenesis and perinatal outcomes: report of the Quality Standards Subcommittee and Therapeutics and Technology Assessment Subcommittee of the American Academy of Neurology and American Epilepsy Society. *Neurology*. 2009;73(2):133–141.
15. Tomson T, Marson A, Boon P, Canevini MP, Covanis A, Gaily E, Kälviäinen R, Trinka E. Valproate in the treatment of epilepsy in girls and women of childbearing potential. *Epilepsia*. 2015;56(7):1006-19.
  16. Chen YH, Chiou HY, Lin HC, Lin HL. Affect of seizures during gestation on pregnancy outcomes in women with epilepsy. *Arch Neurol*. 2009;66(8):979-84.
  17. Rauchenzauner M, Ehrensberger M, Prieschl M, Kapelari K, Bergmann M, Walser G, Neururer S, Unterberger I, Luef G. Generalized tonic-clonic seizures and antiepileptic drugs during pregnancy--a matter of importance for the baby? *J Neurol*. 2013;260(2):484-8.
  18. Veiby G, Daltveit AK, Engelsen BA, Gilhus NE. Pregnancy, delivery, and outcome for the child in maternal epilepsy. *Epilepsia*. 2009;50(9):2130-9.
  19. Man SL, Petersen I, Thompson M, Nazareth I. Antiepileptic drugs during pregnancy in primary care: a UK population based study. *PLoS One*. 2012;7(12):e52339.
  20. Thomas RH, Berkovic SF. The hidden genetics of epilepsy-a clinically important new paradigm. *Nat Rev Neurol*. 2014;10(5):283-92.
  21. Epi4K consortium; Epilepsy Phenome/Genome Project. Ultra-rare genetic variation in common epilepsies: a case-control sequencing study. *Lancet Neurol*. 2017;16(2):135-143.
  22. Viale L, Allotey J, Cheong-See F, et al. Epilepsy in pregnancy and reproductive outcomes: a systematic review and meta-analysis. *Lancet*. 2015;386(10006):1845-52.
  23. Perucca P, O'Brien TJ, Eadie M, Vajda FJ. Valproate still has a place in women with epilepsy. *Epilepsia*. 2015;56(7):1175-6.
  24. Pickrell WO, Lacey AS, Thomas RH, Lyons RA, Smith PE, Rees MI. Trends in the first antiepileptic drug prescribed for epilepsy between 2000 and 2010. *Seizure*. 2014;23(1):77-80.
  25. Pickrell WO, Lacey AS, Bodger OG, Demmler JC, Thomas RH, Lyons RA, Smith PEM, Rees MI, Kerr MP. Epilepsy and deprivation, a data linkage study. *Epilepsia* 2015; 56 (4), 585-591.
  26. Marson AG, Al-Kharusi AM, Alwaidh M, et al. The SANAD study of effectiveness of valproate, lamotrigine, or topiramate for generalised and unclassifiable epilepsy: an unblinded randomised controlled trial. *Lancet*. 2007;369(9566):1016-26.
  27. Edey S, Moran N, Nashef L. SUDEP and epilepsy-related mortality in pregnancy. *Epilepsia*. 2014;55(7):e72-4.
  28. Jones LA, Thomas RH. Sudden death in epilepsy: Insights from the last 25 years *Seizure* 2017;44:232–236.
  29. Kapoor D, Wallace S. Trends in maternal deaths from epilepsy in the United Kingdom: a 30-year retrospective review. *Obstet Med*. 2014;7(4):160-4.
  30. Perucca E. Pharmacological and therapeutic properties of valproate: a summary after 35 years of clinical experience *CNS Drugs* 200;16(10):695–714.
  31. Löscher W. The discovery of valproate. In: Löscher W, editor. *Valproate*. Basel: Birkhäuser, 1999: 1–3.
  32. Mabunga DF, Gonzales EL, Kim JW, Kim KC, Shin CY. Exploring the Validity of Valproic Acid Animal Model of Autism. *Exp Neurobiol*. 2015;24(4):285-300.

33. Martin HG, Manzoni OJ. Late onset deficits in synaptic plasticity in the valproic acid rat model of autism. *Front Cell Neurosci.* 2014;8:23.
34. Walsh J, Thomas RH, Church C, Rees MI, Marson AG, Baker GA. Executive functions and psychiatric symptoms in drug-refractory juvenile myoclonic epilepsy. *Epilepsy Behav.* 2014;35:72-7.
35. Thomas RH, Walsh J, Church C, Sills GJ, Marson AG, Baker GA, Rees MI. A comprehensive neuropsychological description of cognition in drug-refractory juvenile myoclonic epilepsy. *Epilepsy Behav.* 2014;36:124-9.
36. Janz, D and Fuchs, U. Are antiepileptic drugs harmful when given during pregnancy? *Ger Med Mon.* 1964;9:20–22.
37. Bromfield EB, Dworetzky BA, Wyszynski DF, Smith CR, Baldwin EJ, Holmes LB. Valproate teratogenicity and epilepsy syndrome. *Epilepsia.* 2008;49(12):2122-4.
38. Thomas RH, Smith PEM. Epilepsy is different. *Journal of the Royal Society of Medicine* 2011;104(4):141-143.
39. Anazi S, Maddirevula S, Faqeih E, et al. Clinical genomics expands the morbid genome of intellectual disability and offers a high diagnostic yield. *Mol Psychiatry.* 2017;22(4):615-624.
40. Wright CF, Fitzgerald TW, Jones WD, et al. Genetic diagnosis of developmental disorders in the DDD study: a scalable analysis of genome-wide research data. *Lancet.* 2015;385(9975):1305-14.
41. Pennings JL, van Dartel DA, Robinson JF, Pronk TE, Piersma AH. Gene set assembly for quantitative prediction of developmental toxicity in the embryonic stem cell test. *Toxicology.* 2011;284(1-3):63-71.
42. Thomas RH, Robertson NP. The consequences of valproate exposure in utero. *J Neurol.* 2016;263(9):1887-9.
43. [www.OpenPrescribing.net](http://www.OpenPrescribing.net) EBM DataLab, University of Oxford, 2017